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July 24, 2017

Document Processing Desk (ESUB)
Office of Pesticide Programs
U.S. Environmental Protection Agency
One Potomac Yard
2777 South Crystal Drive, Room S4900
Arlington, VA 22202-4501

Attention: Ms Khue Nguyen

Chemical Review Manager

Pesticide Re-evaluation Division (PRD)

Subject: Glyphosate, EPA Reg. No. 524-420;

Glyphosate Technical, EPA Reg. No. 524-421;

Submission of Data for Registration Review of Glyphosate

Dear Ms. Nguyen.

Monsanto herein submits the following data that does not fulfill any specific data requirement other than to support the continued registration of glyphosate herbicide in the U.S.

- Monsanto Glyphosate Manufacturing Industrial Hygiene Monitoring Data, Luling, Louisiana, USA
- Mortality Surveillance Results for the Luling Plant 1980-89
- List of relevant studies to evaluate immunotoxicity
- DRAFT Lymphoma risk and pesticide use in the Agricultural Health Study, Alavanja MCR, DrPH, et al., printed 12/05/2016, produced to Monsanto during personal injury litigation
- 03/20/17 Deposition of Aaron E. Blair, PhD, a listed author of the above DRAFT Report, which
 provides context for the report's findings reglyphosate (taken in above-noted personal injury
 litigation, with p 82-85 removed due to confidentiality provisions)

Since this information is being submitted for overall toxicological considerations of glyphosate as part of the Registration Review of Glyphosate, no PRIA service fee is required.

If you should have any questions regarding this submission, please contact me directly at (314) 694-8270, or by electronic mail at milica vogt@monsanto com, or Dr. Amelia Jackson-Gheissari in the Monsanto D.C. office at (202) 383-2847.

Sincerely,

Milica Vogt

Regulatory Affairs Manager

Milica Vogt

cc⁻ Mr. Reuben Baris / Product Manager, Registration Division, Herbicide Branch Team 25 Dr Amelia Jackson-Gheissari / Monsanto DC Office

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